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(54) Title: LAUNDRY DETERGENT COMPOSITIONS COMPRISING DYE FIXATIVES

(57) Abstract

Disclosed are detergent compositions and methods which utilize certain dye fixatives as fabric treatment agents that can impart fabric appearance benefits to fabrics and textiles laundered in washing solutions which contain such agents. Such dye fixatives are those which do not precipitate with anionic surfactants present in the detergent compositions and which do not therefore adversely affect the cleaning performance of such detergent products.

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LAUNDRY DETERGENT COMPOSITIONS COMPRISING DYE FIXATIVES

TECHNICAL FIELD

The present invention relates to heavy duty laundry detergent compositions, in either liquid or granular form, which contain certain types of dye fixative materials to impart appearance benefits to fabrics and textiles laundered in washing solutions formed from such compositions.

BACKGROUND OF THE INVENTION

It is, of course, well known that alternating cycles of using and laundering fabrics and textiles, such as articles of worn clothing and apparel, will inevitably adversely affect the appearance of the fabric and textile items so used and laundered. Fabrics and textiles simply wear out over time and with use. Laundering of fabrics and textiles is necessary to remove soils and stains which accumulate therein and thereon during ordinary use. However, the laundering operation itself, over many cycles, can accentuate and contribute to the deterioration of the appearance of such fabrics and textiles.

Repeated laundering of fabrics and textiles, especially with bleach-containing laundry products, can remove dye from fabrics and textiles and impart a faded, worn out appearance as a result of diminished color intensity, and in many cases, as a result of changes in hues or shades of color. Dye fixative materials have been used in the textile indusrty to improve the washfastness of certain dyes. Typically such materials are cationic polymers which are able to form ion pairs with the dye on fabrics, thereby reducing the solubility of the dye. Such cationic dye fixative materials, however, cannot typically be used in laundry detergents due to the potential of the cationic

fixative to form precipitates with the anionic surfactants that are generally employed in laundry detergent products. Such precipitation can reduce the cleaning efficiency of the detergent.

Given the foregoing, there is clearly an ongoing need to identify materials which could be added to anionic surfactant-containing laundry detergent products and which could associate themselves with the fibers of the fabrics and textiles laundered using such detergent products and thereby reduce or minimize the tendency of the laundered fabric/textiles to deteriorate in appearance. Any such detergent product additive material should, of course, be able to benefit fabric appearance without unduly interfering with the ability of the laundry detergent to perform its fabric cleaning function. The present invention is directed to detergent compositions containing certain types of dye fixative materials that perform in this desired manner.

SUMMARY OF THE INVENTION

The laundry detergent compositions herein comprise from about 1% to 80% by weight of a detersive surfactant which comprises an anionic surfactant component, from about 0.1% to 80% by weight of an organic or inorganic detergency builder, and from about 0.1% to 5% by weight of certain types of non-precipitating dye fixative materials. The detersive surfactant and detergency builder materials can be any of those useful in conventional laundry detergent products. The useful dye fixative materials are selected from those marketed under the tradenames Sandofix SWE®, Sandofix WA®, Cassofix FRN-300®, Tinofix EW®, Sandolec CT®, Sandolec CS®, Sandolec C1®, Sandolec CF®, Sandolec WA® and Polymer VRN®. Combinations of these selected dye fixative materials may also be used.

In its method aspect, the present invention relates to the laundering of fabrics and textiles in aqueous washing solutions formed from effective amounts of the detergent compositions described herein, or formed from the individual components of such compositions. Laundering of fabrics and textiles in such washing solutions, followed by rinsing and drying, imparts fabric appearance benefits to the fabric and textile articles so treated. Such benefits can include improved overall appearance, and antifading.

DETAILED DESCRIPTION OF THE INVENTION

As noted, the laundry detergent compositions of the present invention essentially contain anionic detersive surfactant, detergent builder and certain selected non-precipitating dye fixative materials which serve to enhance fabric appearance upon use of the detergent compositions to launder fabrics and textiles. Each of these essential

detergent composition components, as well as optional ingredients for such compositions and methods of using such compositions, are described in detail as follows: All percentages and ratios given are by weight unless other specified.

A) Anionic-Containing Detersive Surfactant

The detergent compositions herein essentially comprise from about 1% to 80% by weight of a detersive surfactant which contains an anionic surfactant component. Preferably such compositions comprise from about 5% to 50% by weight of this surfactant. Anionic surfactants can be utilized in the detersive surfactant component in combination with surfactants of the nonionic, zwitterionic, ampholytic or cationic types and will preferably comprise compatible mixtures of these types. Detergent surfactants useful herein are described in U.S. Patent 3,664,961, Norris, Issued May 23, 1972; U.S. Patent 3,919,678, Laughlin et al., Issued December 30, 1975; U.S. Patent 4,222,905, Cockrell, Issued September 16, 1980; and in U.S. Patent 4,239,659, Murphy, Issued December 16, 1980. All of these patents are incorporated herein by reference. Of all the surfactants, combinations of anionics and nonionics are preferred.

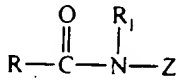
Useful anionic surfactants can themselves be of several different types. For example, water-soluble salts of the higher fatty acids, i.e., "soaps", are useful anionic surfactants in the compositions herein. This includes alkali metal soaps such as the sodium, potassium, ammonium, and alkylolammonium salts of higher fatty acids containing from about 8 to about 24 carbon atoms, and preferably from about 12 to about 18 carbon atoms. Soaps can be made by direct saponification of fats and oils or by the neutralization of free fatty acids. Particularly useful are the sodium and potassium salts of the mixtures of fatty acids derived from coconut oil and tallow, i.e., sodium or potassium tallow and coconut soap.

Additional non-soap anionic surfactants which are suitable for use herein include the water-soluble salts, preferably the alkali metal, and ammonium salts, of organic sulfuric reaction products having in their molecular structure an alkyl group containing from about 10 to about 20 carbon atoms and a sulfonic acid or sulfuric acid ester group. (Included in the term "alkyl" is the alkyl portion of acyl groups.) Examples of this group of synthetic surfactants are a) the sodium, potassium and ammonium alkyl sulfates, especially those obtained by sulfating the higher alcohols (C_8-C_{18} carbon atoms) such as those produced by reducing the glycerides of tallow or coconut oil; b) the sodium, potassium and ammonium alkyl polyethoxylate sulfates, particularly those in which the alkyl group contains from 10 to 22, preferably from 12 to 18 carbon atoms, and wherein the polyethoxylate chain contains from 1 to 15, preferably 1 to 6 ethoxylate moieties; and c) the sodium and potassium alkylbenzene sulfonates in which

the alkyl group contains from about 9 to about 15 carbon atoms, in straight chain or branched chain configuration, e.g., those of the type described in U.S. Patents 2,220,099 and 2,477,383. Especially valuable are linear straight chain alkylbenzene sulfonates in which the average number of carbon atoms in the alkyl group is from about 11 to 13, abbreviated as C_{11-13} LAS.

Preferred nonionic surfactants for use in combination with the foregoing anionics are those of the formula $R^1(OC_2H_4)_nOH$, wherein R^1 is a C_{10}-C_{16} alkyl group or a C_8-C_{12} alkylphenol group, and n is from 3 to about 80. Particularly preferred are condensation products of C_{12}-C_{15} alcohols with from about 5 to about 20 moles of ethylene oxide per mole of alcohol, e.g., C_{12}-C_{13} alcohol condensed with about 6.5 moles of ethylene oxide per mole of alcohol.

Additional suitable nonionic surfactants include polyhydroxy fatty acid amides of the formula:



wherein R is a C_{9-17} alkyl or alkenyl, R_1 is a methyl group and Z is glycityl derived from a reduced sugar or alkoxylated derivative thereof. Examples are N-methyl N-1-deoxyglucityl cocoamide and N-methyl N-1-deoxyglucityl oleamide. Processes for making polyhydroxy fatty acid amides are known and can be found in Wilson, U.S. Patent 2,965,576 and Schwartz, U.S. Patent 2,703,798, the disclosures of which are incorporated herein by reference.

B) Detergent Builder

The detergent compositions herein also essentially comprise from about 0.1% to 80% by weight of a detergent builder. Preferably such compositions in liquid form will comprise from about 1% to 10% by weight of the builder component. Preferably such compositions in granular form will comprise from about 1% to 50% by weight of the builder component. Detergent builders are well known in the art and can comprise, for example, phosphate salts as well as various organic and inorganic nonphosphorus builders.

Water-soluble, nonphosphorus organic builders useful herein include the various alkali metal, ammonium and substituted ammonium polyacetates, carboxylates, polycarboxylates and polyhydroxy sulfonates. Examples of polyacetate and polycarboxylate builders are the sodium, potassium, lithium, ammonium and substituted ammonium salts of ethylene diamine tetraacetic acid, nitrilotriacetic acid, oxydisuccinic acid, mellitic acid, benzene polycarboxylic acids, and citric acid. Other

suitable polycarboxylates for use herein are the polyacetal carboxylates described in U.S. Patent 4,144,226, issued March 13, 1979 to Crutchfield et al, and U.S. Patent 4,246,495, issued March 27, 1979 to Crutchfield et al, both of which are incorporated herein by reference. Particularly preferred polycarboxylate builders are the oxydisuccinates and the ether carboxylate builder compositions comprising a combination of tartrate monosuccinate and tartrate disuccinate described in U.S. Patent 4,663,071, Bush et al., issued May 5, 1987, the disclosure of which is incorporated herein by reference.

Examples of suitable nonphosphorus, inorganic builders include the silicates, aluminosilicates, borates and carbonates. Particularly preferred are sodium and potassium carbonate, bicarbonate, sesquicarbonate, tetraborate decahydrate, and silicates having a weight ratio of SiO_2 to alkali metal oxide of from about 0.5 to about 4.0, preferably from about 1.0 to about 2.4. Also preferred are aluminosilicates including zeolites. Such materials and their use as detergent builders are more fully discussed in Corkill et al, U. S. Patent No. 4,605,509, the disclosure of which is incorporated herein by reference. Also, crystalline layered silicates such as those discussed in Corkill et al, U. S. Patent No. 4,605,509, incorporated herein by reference, are suitable for use in the detergent compositions of this invention.

C) Dye Fixative Materials

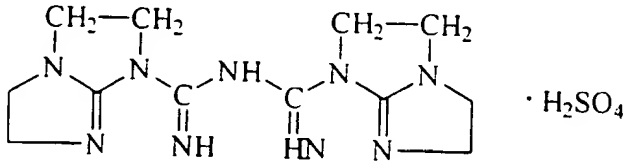
The third essential component of the detergent compositions herein comprises selected dye fixative materials which do not form precipitates with the essentially-utilized anionic surfactant. Such non-precipitating dye fixative materials have been found to impart appearance benefits to fabrics and textiles laundered in aqueous washing solutions formed from detergent compositions which contain such dye fixatives. These fabric appearance benefits can include, for example, improved overall appearance of the laundered fabrics and protection against color fading. The selected dye fixatives used in the compositions and methods herein can provide such fabric appearance benefits with acceptably little or no loss in cleaning performance provided by the laundry detergent compositions into which such materials are incorporated.

The selected dye fixatives useful herein may be in the form of unpolymerized materials, oligomers or polymers. The dye fixative component of the compositions herein will generally comprise from about 0.1% to 5% by the weight of the composition. More preferably, such dye fixative materials will comprise from about 0.5% to 4% by weight of the compositions, most preferably from about 1% to 3%. Such concentrations should be sufficient to provide from about 10 to 100 ppm of the dye fixative in the aqueous washing solutions formed from the laundry detergent

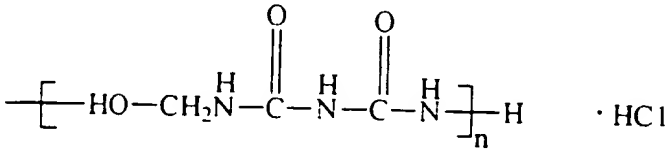
compositions herein. More prefearably from about 20 to 60 ppm of the dye fixative will be delivered to the aqueous washing solution, most preferably about 50 ppm.

The non-precipitating dye fixatives useful herein include a number that are commercially marketed by CLARIANT Corporation under the Sandofix®, Sandolec® and Polymer VRN® tradenames. These include, for example, Sandofix SWE®, Sandofix WA®, Sandolec CT®, Sandolec CS®, Sandolec C1®, Sandolec CF®, Sandolec WA® and Polymer VRN®. Other suitable dye fixatives are marketed by Ciba-Geigy Corporation under the tradename Tinofix EW® and by Hoechst Celanese Corporation under the tradename Cassofix FRN-300®.

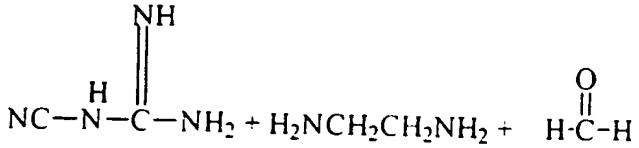
Preferred non-precipitating dye fixative materials are Sandofix SWE® and Sandolec CS® which have the structure:



Other preferred non-precipitating dye fixative materials are Sandofix WA® and Sandolec WA® which are polymers having the following structure:

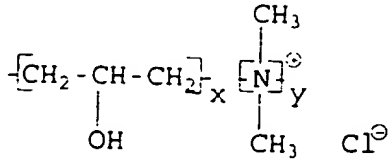


Another preferred non-precipitating dye fixative is the Cassofix FRN-300® material which is also a polymer prepared from the following monomers:

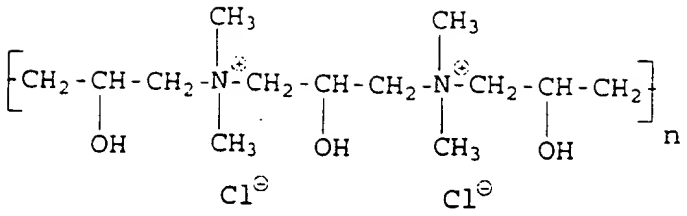


Chemical structures for some additionally preferred dye fixative compounds for use in the present invention are as follows:

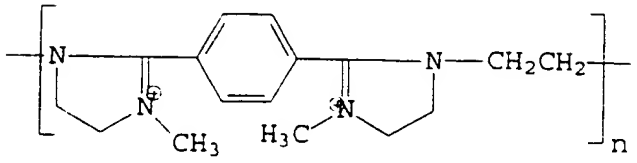
Sandolec CT®,



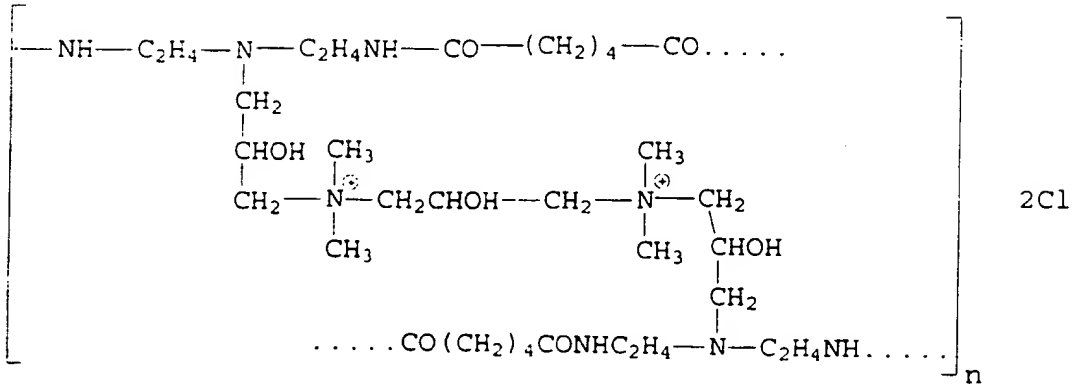
Polymer VRN®,



Sandolec Cl®,



Sandolec CF®,



The dye fixative materials used in this invention are generally all water-soluble materials. They can therefore be utilized for detergent composition preparation in the form of aqueous solutions of such dye fixatives if desired.

D) Optional Detergent Ingredients

In addition to the essential surfactants, builders and dye fixatives hereinbefore described, the detergent composition of the present invention can also include any number of additional optional ingredients. These include conventional detergent composition components such as bleaches and bleach activators, enzymes and enzyme stabilizing agents, suds boosters or suds suppressers, anti-tarnish and anticorrosion agents, soil suspending agents, soil release agents, germicides, pH adjusting agents, non-builder alkalinity sources, chelating agents, organic and inorganic fillers, solvents, hydrotropes, optical brighteners, dyes and perfumes.

A preferred optional ingredients for incorporation into the detergent compositions herein comprises a bleaching agent, e.g., a peroxygen bleach. Such peroxygen bleaching agents may be organic or inorganic in nature. Inorganic peroxygen bleaching agents are frequently utilized in combination with a bleach activator.

Useful organic peroxygen bleaching agents include percarboxylic acid bleaching agents and salts thereof. Suitable examples of this class of agents include magnesium monoperoxyphthalate hexahydrate, the magnesium salt of metachloro perbenzoic acid, 4-nonylamino-4-oxoperoxybutyric acid and diperoxydodecanedioic acid. Such bleaching agents are disclosed in U.S. Patent 4,483,781, Hartman, Issued November 20, 1984; European Patent Application EP-A-133,354, Banks et al., Published February 20, 1985; and U.S. Patent 4,412,934, Chung et al., Issued November 1, 1983. Highly preferred bleaching agents also include 6-nonylamino-6-oxoperoxycaproic acid (NAPAA) as described in U.S. Patent 4,634,551, Issued January 6, 1987 to Burns et al.

Inorganic peroxygen bleaching agents may also be used, generally in particulate form, in the detergent compositions herein. Inorganic bleaching agents are in fact preferred. Such inorganic peroxygen compounds include alkali metal perborate and percarbonate materials. For example, sodium perborate (e.g. mono- or tetra-hydrate) can be used. Suitable inorganic bleaching agents can also include sodium or potassium carbonate peroxyhydrate and equivalent "percarbonate" bleaches, sodium pyrophosphate peroxyhydrate, urea peroxyhydrate, and sodium peroxide. Persulfate bleach (e.g., OXONE, manufactured commercially by DuPont) can also be used. Frequently inorganic peroxygen bleaches will be coated with silicate, borate, sulfate or water-soluble surfactants. For example, coated percarbonate particles are available from various commercial sources such as FMC, Solvay Interox, Tokai Denka and Degussa.

Inorganic peroxygen bleaching agents, e.g., the perborates, the percarbonates, etc., are preferably combined with bleach activators, which lead to the *in situ* production in aqueous solution (i.e., during use of the compositions herein for fabric laundering/bleaching) of the peroxy acid corresponding to the bleach activator. Various non-limiting examples of activators are disclosed in U.S. Patent 4,915,854, Issued April 10, 1990 to Mao et al.; and U.S. Patent 4,412,934 Issued November 1, 1983 to Chung et al. The nonanoyloxybenzene sulfonate (NOBS) and tetraacetyl ethylene diamine (TAED) activators are typical and preferred. Mixtures thereof can also be used. See also the hereinbefore referenced U.S. 4,634,551 for other typical bleaches and activators useful herein.

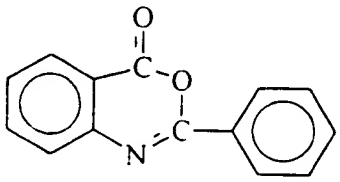
Other useful amido-derived bleach activators are those of the formulae:

$$R^1N(R^5)C(O)R^2C(O)L \quad \text{or} \quad R^1C(O)N(R^5)R^2C(O)L$$

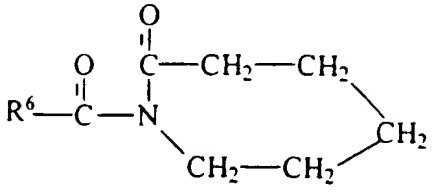
wherein R^1 is an alkyl group containing from about 6 to about 12 carbon atoms, R^2 is an alkylene containing from 1 to about 6 carbon atoms, R^5 is H or alkyl, aryl, or alkaryl containing from about 1 to about 10 carbon atoms, and L is any suitable leaving group. A leaving group is any group that is displaced from the bleach activator as a consequence of the nucleophilic attack on the bleach activator by the perhydrolysis anion. A preferred leaving group is phenol sulfonate.

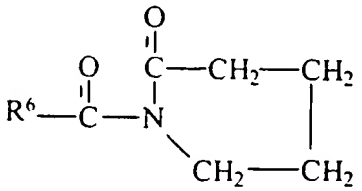
Preferred examples of bleach activators of the above formulae include (6-octanamido-caproyl)oxybenzenesulfonate, (6-nonanamidocaproyl) oxybenzenesulfonate, (6-decanamido-caproyl)oxybenzenesulfonate and mixtures thereof as described in the hereinbefore referenced U.S. Patent 4,634,551.

Another class of useful bleach activators comprises the benzoxazin-type activators disclosed by Hodge et al. in U.S. Patent 4,966, 723, Issued October 30, 1990, incorporated herein by reference. A highly preferred activator of the benzoxazin-type is:



Still another class of useful bleach activators includes the acyl lactam activators, especially acyl caprolactams and acyl valerolactams of the formulae:





wherein R^6 is H or an alkyl, aryl, alkoxyaryl, or alkaryl group containing from 1 to about 12 carbon atoms. Highly preferred lactam activators include benzoyl caprolactam, octanoyl caprolactam, 3,5,5-trimethylhexanoyl caprolactam, nonanoyl caprolactam, decanoyl caprolactam, undecenoyl caprolactam, benzoyl valerolactam, octanoyl valerolactam, nonanoyl valerolactam, decanoyl valerolactam, undecenoyl valerolactam, 3,5,5-trimethylhexanoyl valerolactam and mixtures thereof. See also U.S. Patent 4,545,784, Issued to Sanderson, October 8, 1985, incorporated herein by

reference, which discloses acyl caprolactams, including benzoyl caprolactam, adsorbed into sodium perborate.

If utilized, peroxygen bleaching agent will generally comprise from about 2% to 30% by weight of the detergent compositions herein. More preferably, peroxygen bleaching agent will comprise from about 2% to 20% by weight of the compositions. Most preferably, peroxygen bleaching agent will be present to the extent of from about 3% to 15% by weight of the compositions herein. If utilized, bleach activators can comprise from about 2% to 10% by weight of the detergent compositions herein. Frequently, activators are employed such that the molar ratio of bleaching agent to activator ranges from about 1:1 to 10:1, more preferably from about 1.5:1 to 5:1.

Another highly preferred optional ingredient in the detergent compositions herein is a detersive enzymes component. Enzymes can be included in the present detergent compositions for a variety of purposes, including removal of protein-based, carbohydrate-based, or triglyceride-based stains from substrates, for the prevention of refugee dye transfer in fabric laundering, and for fabric restoration. Suitable enzymes include proteases, amylases, lipases, cellulases, peroxidases, and mixtures thereof of any suitable origin, such as vegetable, animal, bacterial, fungal and yeast origin. Preferred selections are influenced by factors such as pH-activity and/or stability optima, thermostability, and stability to active detergents, builders and the like. In this respect bacterial or fungal enzymes are preferred, such as bacterial amylases and proteases, and fungal cellulases.

"Detersive enzyme", as used herein, means any enzyme having a cleaning, stain removing or otherwise beneficial effect in a laundry detergent composition. Preferred enzymes for laundry purposes include, but are not limited to, proteases, cellulases, lipases, amylases and peroxidases.

Enzymes are normally incorporated into detergent compositions at levels sufficient to provide a "cleaning-effective amount". The term "cleaning-effective amount" refers to any amount capable of producing a cleaning, stain removal, soil removal, whitening, deodorizing, or freshness improving effect on substrates such as fabrics. In practical terms for current commercial preparations, typical amounts are up to about 5 mg by weight, more typically 0.01 mg to 3 mg, of active enzyme per gram of the detergent composition. Stated otherwise, the compositions herein will typically comprise from 0.001% to 5%, preferably 0.01%-1% by weight of a commercial enzyme preparation. Protease enzymes are usually present in such commercial preparations at levels sufficient to provide from 0.005 to 0.1 Anson units (AU) of activity per gram of

composition. Higher active levels may be desirable in highly concentrated detergent formulations.

Suitable examples of proteases are the subtilisins which are obtained from particular strains of *B. subtilis* and *B. licheniformis*. One suitable protease is obtained from a strain of *Bacillus*, having maximum activity throughout the pH range of 8-12, developed and sold as ESPERASE® by Novo Industries A/S of Denmark, hereinafter "Novo". The preparation of this enzyme and analogous enzymes is described in GB 1,243,784 to Novo. Other suitable proteases include ALCALASE® and SAVINASE® from Novo and MAXATASE® from International Bio-Synthetics, Inc., The Netherlands; as well as Protease A as disclosed in EP 130,756 A, January 9, 1985 and Protease B as disclosed in EP 303,761 A, April 28, 1987 and EP 130,756 A, January 9, 1985. See also a high pH protease from Bacillus sp. NCIMB 40338 described in WO 9318140 A to Novo. Enzymatic detergents comprising protease, one or more other enzymes, and a reversible protease inhibitor are described in WO 9203529 A to Novo. Other preferred proteases include those of WO 9510591 A to Procter & Gamble . When desired, a protease having decreased adsorption and increased hydrolysis is available as described in WO 9507791 to Procter & Gamble. A recombinant trypsin-like protease for detergents suitable herein is described in WO 9425583 to Novo.

Cellulases usable herein include both bacterial and fungal types, preferably having a pH optimum between 5 and 10. U.S. 4,435,307, Barbesgoard et al, March 6, 1984, discloses suitable fungal cellulases from *Humicola insolens* or *Humicola* strain DSM1800 or a cellulase 212-producing fungus belonging to the genus *Aeromonas*, and cellulase extracted from the hepatopancreas of a marine mollusk, *Dolabella Auricula Solander*. Suitable cellulases are also disclosed in GB-A-2,075,028; GB-A-2,095,275 and DE-OS-2,247,832. CAREZYME® and CELLUZYME® (Novo) are especially useful. See also WO 9117243 to Novo.

Suitable lipase enzymes for detergent usage include those produced by microorganisms of the *Pseudomonas* group, such as *Pseudomonas stutzeri* ATCC 19.154, as disclosed in GB 1,372,034. See also lipases in Japanese Patent Application 53,20487, laid open Feb. 24, 1978. This lipase is available from Amano Pharmaceutical Co. Ltd., Nagoya, Japan, under the trade name Lipase P "Amano," or "Amano-P." Other suitable commercial lipases include Amano-CES, lipases ex *Chromobacter viscosum*, e.g. *Chromobacter viscosum var. lipolyticum* NRRLB 3673 from Toyo Jozo Co., Tagata, Japan; *Chromobacter viscosum* lipases from U.S. Biochemical Corp., U.S.A. and Disoynth Co., The Netherlands, and lipases ex *Pseudomonas gladioli*. LIPOLASE® enzyme derived from *Humicola lanuginosa* and

commercially available from Novo, see also EP 341,947, is a preferred lipase for use herein.

The enzyme-containing compositions herein may optionally also comprise from about 0.001% to about 10%, preferably from about 0.005% to about 8%, most preferably from about 0.01% to about 6%, by weight of an enzyme stabilizing system. The enzyme stabilizing system can be any stabilizing system which is compatible with the detersive enzyme. Such a system may be inherently provided by other formulation actives, or be added separately, e.g., by the formulator or by a manufacturer of detergent-ready enzymes. Such stabilizing systems can, for example, comprise calcium ion, boric acid, propylene glycol, short chain carboxylic acids, boronic acids, and mixtures thereof, and are designed to address different stabilization problems depending on the type and physical form of the detergent composition.

E) Detergent Composition Preparation

The detergent compositions according to the present invention can be in liquid, paste or granular forms. Such compositions can be prepared by combining the essential and optional components in the requisite concentrations in any suitable order and by any conventional means.

Granular compositions, for example, are generally made by combining base granule ingredients (e.g. surfactants, builders, water, etc.) as a slurry, and spray drying the resulting slurry to a low level of residual moisture (5-12%). The remaining dry ingredients can be admixed in granular powder form with the spray dried granules in a rotary mixing drum and the liquid ingredients (e.g. aqueous solutions of the essential dye fixatives, enzymes, binders and perfumes) can be sprayed onto the resulting granules to form the finished detergent composition. Granular compositions according to the present invention can also be in "compact form", i.e. they may have a relatively higher density than conventional granular detergents, i.e. from 550 to 950 g/l. In such case, the granular detergent compositions according to the present invention will contain a lower amount of "inorganic filler salt", compared to conventional granular detergents; typical filler salts are alkaline earth metal salts of sulphates and chlorides, typically sodium sulphate; "compact" detergents typically comprise not more than 10% filler salt.

Liquid detergent compositions can be prepared by admixing the essential and optional ingredients thereof in any desired order to provide compositions containing components in the requisite concentrations. Liquid compositions according to the

present invention can also be in "compact form", in such case, the liquid detergent compositions according to the present invention will contain a lower amount of water, compared to conventional liquid detergents.

Addition of the dye fixative component to liquid detergent compositions of this invention may be accomplished by simply mixing into the liquid dertergent aqueous solutions of the desired dye fixatives. Such dye fixatives can alter the viscosity or other rheological characteristics of liquid detergent products. It may therefore be necessary to compensate for any rheological changes in the liquid detergent product brought about by dye fixative addition by altering the type and amount of hydrotropes and/or solvents that are used.

F) Fabric Laundering Method

The present invention also provides a method for laundering fabrics in a manner which imparts fabric appearance benefits provided by the dye fixative materials used herein. Such a method employs contacting these fabrics with an aqueous washing solution formed from an effective amount of the detergent compositions hereinbefore described or formed from the individual components of such compositions. Contacting of fabrics with washing solution will generally occur under conditions of agitation although the compositions of the present invention may also be used to form aqueous unagitated soaking solutions for fabric cleaning and treatment.

Agitation is preferably provided in a washing machine for good cleaning. Washing is preferably followed by drying the wet fabric in a conventional clothes dryer. An effective amount of the liquid or granular detergent composition in the aqueous wash solution in the washing machine is preferably from about 500 to about 7000 ppm, more preferably from about 1000 to about 3000 ppm.

EXAMPLES

The following examples illustrate the compositions and methods of the present invention, but are not necessarily meant to limit or otherwise define the scope of the invention.

EXAMPLES 1-33

Liquid Detergent Test Composition Preparation

Several heavy duty liquid (HDL) detergent compositions are prepared containing various dye fixatives. Such liquid detergent compositions all have the following basic formula:

Component	Wt. %
C_{12-15} alkyl ether (2.5) sulfate	19.0
C_{12-13} alkyl ethoxylate (9.0)	2.00
C_{12-14} glucose amide	3.50
Citric Acid	3.00
C_{12-14} Fatty Acid	2.00
MEA	to pH 8
Ethanol	3.41
Propanediol	6.51
Borax	2.5
Dispersant	1.18
Na Toluene Sulfonate	2.50
Dye Fixative as shown in Table 1	See Table 1
Dye, Perfume, Brighteners, Enzymes, Preservatives, Suds Suppressor, Other Minors, Water	Balance
	100%

Table 1. Dye Fixatives Used in Liquid Test Detergents

Example #	Dye Fixative	Supplier	Wt. % Active in HDL
1	Sandofix SWE	Clariant	4.8
2	Sandofix SWE	Clariant	3.2
3	Sandofix SWE	Clariant	2.0
4	Sandofix SWE	Clariant	1.6
5	Sandofix SWE	Clariant	1.3
6	Sandofix SWE	Clariant	0.6
7	Cassofix FRN-300	Hoechst	4.8
8	Cassofix FRN-300	Hoechst	3.2
9	Cassofix FRN-300	Hoechst	1.6
10	Cassofix FRN-300	Hoechst	1.3
11	Cassofix FRN-300	Hoechst	0.6
12	Sandofix WA	Clariant	4.8
13	Sandofix WA	Clariant	3.2
14	Sandofix WA	Clariant	1.6
15	Sandofix WA	Clariant	1.0
16	Sandofix WA	Clariant	0.6
17	Tinofix EW	Ciba	4.8
18	Tinofix EW	Ciba	3.2
19	Tinofix EW	Ciba	1.6
20	Tinofix EW	Ciba	0.6
21	Sandolec CT	Clariant	4.8
22	Sandolec CT	Clariant	3.2
23	Sandolec CT	Clariant	2.0
24	Sandolec CT	Clariant	1.6
25	Sandolec CT	Clariant	0.6
26	Polymer VRN	Clariant	4.8
27	Polymer VRN	Clariant	2.0
28	Sandolec CS	Clariant	4.8
29	Sandolec CS	Clariant	3.2
30	Sandolec Cl	Clariant	4.8
31	Sandolec Cl	Clariant	3.2
32	Sandolec CF	Clariant	4.8

33	Sandolec WA	Clariant	4.8

EXAMPLE 34-36

Granular Detergent Test Composition Preparation

Several heavy duty granular detergent (HDG) compositions are prepared containing various dye fixatives. Such granular detergent compositions all have the following basic formula:

Component	Wt. %
C_{12} Linear alkyl benzene sulfonate	9.31
C_{14-15} alkyl sulfonate	12.74
Zeolite Builder	27.79
Sodium Carbonate	27.31
PEG 4000	1.60
Dispersant	2.26
C_{12-13} alkyl ethoxylate (E9)	1.5
Sodium Perborate	1.03
Soil Release Polymer	0.41
Enzymes	0.59
Dye Fixative	Seen Table 2
Perfume, Brightener, Suds Suppressor, Other Minors, Moisture, Sulfate	Balance
	100%

Table 2. Dye Fixatives Used in Granular Test Detergents

Example #	Polyamide	Supplier	Conc. in Solution
34	Sandofix SWE	Clariant	50 ppm
35	Cassofix FRN-300 (KDM data)	Hoechst	50 ppm
36	Sandofix WA (KDM data)	Clariant	50 ppm

WHAT IS CLAIMED IS:

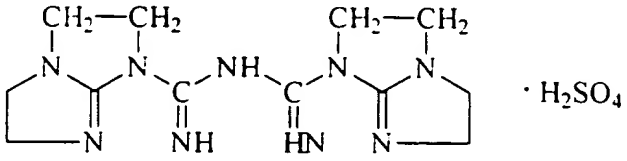
1.A laundry detergent composition which imparts the fabric appearance benefit of antifading to fabrics and textiles laundered in aqueous washing solutions formed therefrom, which composition characterizes:

A) from 5% to 50% by weight of a detersive surfactant component which comprises at least one anionic surfactant;

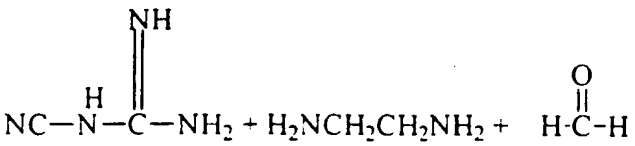
B) from 1% to 50% by weight of an organic or inorganic detergency builder;

C) from 0.1% to 5% by weight of a dye fixative selected from the group consisting of Cassofix FRN-300®, Sandofix SWE®, Sandofix WA®, Tinofix EW®, Polymer VRN®, Sandolec CF®, Sandolec WA®, Sandolec CT®, Sandolec CS® and Sandolec CI®.

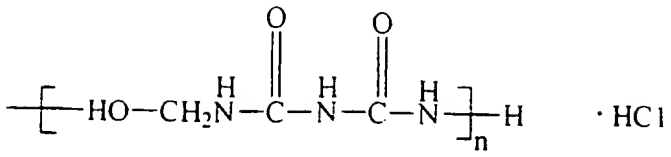
2. A composition according to Claim 1 wherein the dye fixative is Sandofix SWE® having the formula:



3. A compositon according to Claim 1 wherein the dye fixative is a Cassofix FRN-300® polymer prepared from monomers having the following structures:



4. A composition according to Claim 1 wherein the dye fixative is a Sandofix WA® polymer having the following structure:



INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 97/22596

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C11D3/28 C11D3/30 C11D3/37

According to International Patent Classification(IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 C11D D06L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No
A	EP 0 462 806 A (UNILEVER PLC ET AL) 27 December 1991 see page 2, line 39 - page 4, line 33 ---	1
A	WO 96 27649 A (THE PROCTER & GAMBLE COMPANY) 12 September 1996 see page 3, paragraph 1 - page 4, paragraph 3 see page 10, paragraph 2 - page 11, paragraph 2; claims 1,4 ---	1,2
T	WO 97 46650 A (THE PROCTER & GAMBLE COMPANY) 11 December 1997 see page 3, paragraph 4 - page 11, paragraph 2 see page 16, paragraph 5 - page 17, paragraph 2 -----	1,2

☐ Further documents are listed in the continuation of box C

☒ Patent family members are listed in annex

* Special categories of cited documents

"A" document defining the general state of the art which is not considered to be of particular relevance

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"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
19 March 1998	08/04/1998

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Information on patent family members

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PCT/US 97/22596

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